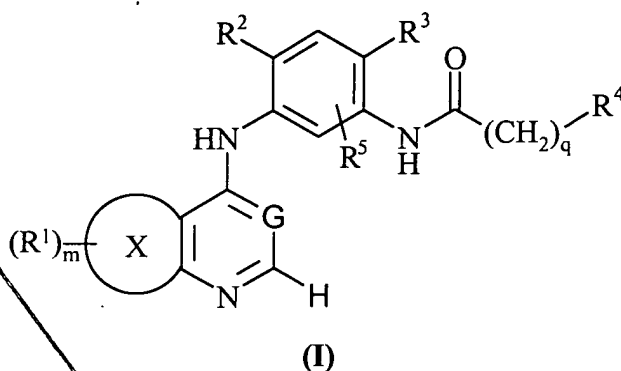


**IN THE CLAIMS:**

Please cancel claim 11, and amend claims 1, 10 and 12 to read as follows, without prejudice to applicant's right to prosecute any cancelled subject matter in one or more divisional or continuing applications:

Sub 1. (Amended) A bicyclic compound of the Formula (I):



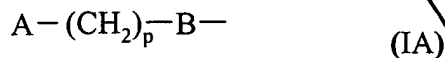
wherein:

G is N;

ring X is a 5- or 6-membered fused heteroaryl ring which contains 1, 2 or 3 heteroatoms selected from oxygen, sulphur and nitrogen;

m is 0, 1 or 2;

R<sup>1</sup> is hydroxy, halo, trifluoromethyl, cyano, mercapto, nitro, amino, carboxy, carbamoyl, formyl, sulphamoyl, C<sub>1-6</sub>alkyl, C<sub>2-6</sub>alkenyl, C<sub>2-6</sub>alkynyl, C<sub>1-6</sub>alkoxy, -O-(C<sub>1-3</sub>alkyl)-O-, C<sub>1-6</sub>alkylS(O)<sub>n</sub>- (wherein n is 0-2), N-C<sub>1-6</sub>alkylamino, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, C<sub>1-6</sub>alkoxycarbonyl, N-C<sub>1-6</sub>alkylcarbamoyl, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, C<sub>2-6</sub>alkanoyl, C<sub>1-6</sub>alkanoyloxy, C<sub>1-6</sub>alkanoylamino, N-C<sub>1-6</sub>alkylsulphamoyl, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>sulphamoyl, C<sub>1-6</sub>alkylsulphonylamino, C<sub>1-6</sub>alkylsulphonyl-N-(C<sub>1-6</sub>alkyl)amino, or R<sup>1</sup> is of the Formula (IA):



wherein A is halo, hydroxy, C<sub>1-6</sub>alkoxy, C<sub>1-6</sub>alkylS(O)<sub>n</sub>- (wherein n is 0-2), cyano, amino, N-C<sub>1-6</sub>alkylamino, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, carboxy, C<sub>1-6</sub>alkoxycarbonyl, carbamoyl, N-C<sub>1-6</sub>alkylcarbamoyl or N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, p is 1 - 6, and B is a bond, oxy, imino, N-(C<sub>1-6</sub>alkyl)imino or -C(O)NH-, with the proviso that p is 2 or more unless B is a bond or -C(O)NH-,

or R<sup>1</sup> is of the Formula (IB):



wherein D is aryl, heteroaryl or heterocyclyl and E is a bond, C<sub>1-6</sub>alkylene, C<sub>1-6</sub>alkyleneoxy, oxy, imino, N-(C<sub>1-6</sub>alkyl)imino, C<sub>1-6</sub>alkyleneimino, N-(C<sub>1-6</sub>alkyl)-C<sub>1-6</sub>alkyleneimino, C<sub>1-6</sub>alkyleneoxy-C<sub>1-6</sub>alkylene, C<sub>1-6</sub>alkyleneimino-C<sub>1-6</sub>alkylene, N-(C<sub>1-6</sub>alkyl)-C<sub>1-6</sub>alkyleneimino-C<sub>1-6</sub>alkylene, -C(O)NH-, -SO<sub>2</sub>NH-, -NHSO<sub>2</sub>- or C<sub>2-6</sub>alkanoylimino,

and any aryl, heteroaryl or heterocyclyl group in a R<sup>1</sup> group may be optionally substituted with one or more groups selected from hydroxy, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, carboxy, C<sub>1-6</sub>alkoxycarbonyl, carbamoyl, N-C<sub>1-6</sub>alkylcarbamoyl, N-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, C<sub>2-6</sub>alkanoyl, amino, N-C<sub>1-6</sub>alkylamino and N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino,

and any heterocyclyl group in a R<sup>1</sup> group may be optionally substituted with one or two oxo or thioxo substituents,

and any of the R<sup>1</sup> groups defined hereinbefore which comprises a CH<sub>2</sub> group which is attached to 2 carbon atoms or a CH<sub>3</sub> group which is attached to a carbon atom may optionally bear on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, amino, C<sub>1-6</sub>alkoxy, N-C<sub>1-6</sub>alkylamino, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino and heterocyclyl;

R<sup>2</sup> is hydrogen, halo, C<sub>1-6</sub>alkyl, C<sub>2-6</sub>alkenyl or C<sub>2-6</sub>alkynyl;

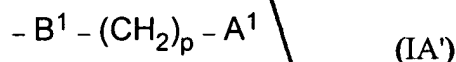
R<sup>3</sup> is hydrogen, halo, C<sub>1-6</sub>alkyl, C<sub>2-6</sub>alkenyl or C<sub>2-6</sub>alkynyl;

R<sup>4</sup> is hydrogen, hydroxy, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, amino, N-C<sub>1-6</sub>alkylamino, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, hydroxyC<sub>2-6</sub>alkoxy, C<sub>1-6</sub>alkoxyC<sub>2-6</sub>alkoxy, aminoC<sub>2-6</sub>alkoxy, N-C<sub>1-6</sub>alkylaminoC<sub>2-6</sub>alkoxy, N,N-(C<sub>1-6</sub>alkyl)<sub>2</sub>aminoC<sub>2-6</sub>alkoxy or C<sub>3-7</sub>cycloalkyl,

or R<sup>4</sup> is of the Formula (IC):



wherein J is aryl, heteroaryl or heterocyclyl and K is a bond, oxy, imino, *N*-(C<sub>1-6</sub>alkyl)imino, oxyC<sub>1-6</sub>alkylene, iminoC<sub>1-6</sub>alkylene, *N*-(C<sub>1-6</sub>alkyl)iminoC<sub>1-6</sub>alkylene, -NHC(O)-, -SO<sub>2</sub>NH-, -NHSO<sub>2</sub>- or -NHC(O)-C<sub>1-6</sub>alkylene-, and any aryl, heteroaryl or heterocyclyl group in a R<sup>4</sup> group may be optionally substituted by one or more groups selected from hydroxy, halo, trifluoromethyl, cyano, mercapto, nitro, amino, carboxy, carbamoyl, formyl, sulphamoyl, C<sub>1-6</sub>alkyl, C<sub>2-6</sub>alkenyl, C<sub>2-6</sub>alkynyl, C<sub>1-6</sub>alkoxy, -O-(C<sub>1-3</sub>alkyl)-O-, C<sub>1-6</sub>alkylS(O)<sub>n</sub>- (wherein n is 0-2), *N*-C<sub>1-6</sub>alkylamino, *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, C<sub>1-6</sub>alkoxycarbonyl, *N*-C<sub>1-6</sub>alkylcarbamoyl, *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, C<sub>2-6</sub>alkanoyl, C<sub>1-6</sub>alkanoyloxy, C<sub>1-6</sub>alkanoylamino, *N*-C<sub>1-6</sub>alkylsulphamoyl, *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>sulphamoyl, C<sub>1-6</sub>alkylsulphonylamino and C<sub>1-6</sub>alkylsulphonyl-*N*-(C<sub>1-6</sub>alkyl)amino, or any aryl, heteroaryl or heterocyclyl group in a R<sup>4</sup> group may be optionally substituted with one or more groups of the Formula (IA'):



wherein A<sup>1</sup> is halo, hydroxy, C<sub>1-6</sub>alkoxy, cyano, amino, *N*-C<sub>1-6</sub>alkylamino, *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, carboxy, C<sub>1-6</sub>alkoxycarbonyl, carbamoyl, *N*-C<sub>1-6</sub>alkylcarbamoyl or *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, p is 1 - 6, and B<sup>1</sup> is a bond, oxy, imino, *N*-(C<sub>1-6</sub>alkyl)imino or -NHC(O)-, with the proviso that p is 2 or more unless B<sup>1</sup> is a bond or -NHC(O)-, or any aryl, heteroaryl or heterocyclyl group in a R<sup>4</sup> group may be optionally substituted with one or more groups of the Formula (IB'):



wherein D<sup>1</sup> is aryl, heteroaryl or heterocyclyl and E<sup>1</sup> is a bond, C<sub>1-6</sub>alkylene, oxyC<sub>1-6</sub>alkylene, oxy, imino, *N*-(C<sub>1-6</sub>alkyl)imino, iminoC<sub>1-6</sub>alkylene, *N*-(C<sub>1-6</sub>alkyl)-iminoC<sub>1-6</sub>alkylene, C<sub>1-6</sub>alkylene-oxyC<sub>1-6</sub>alkylene, C<sub>1-6</sub>alkylene-iminoC<sub>1-6</sub>alkylene, C<sub>1-6</sub>alkylene-*N*-(C<sub>1-6</sub>alkyl)-iminoC<sub>1-6</sub>alkylene, -NHC(O)-, -NHSO<sub>2</sub>-, -SO<sub>2</sub>NH- or -NHC(O)-C<sub>1-6</sub>alkylene-, and any aryl, heteroaryl or heterocyclyl group in a substituent on R<sup>4</sup> may be optionally substituted with one or more groups selected from hydroxy, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy,

B  
contd.

carboxy, C<sub>1-6</sub>alkoxycarbonyl, carbamoyl, *N*-C<sub>1-6</sub>alkylcarbamoyl, *N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>carbamoyl, C<sub>2-6</sub>alkanoyl, amino, *N*-C<sub>1-6</sub>alkylamino and *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino, and any C<sub>3-7</sub>cycloalkyl or heterocyclyl group in a R<sup>4</sup> group may be optionally substituted with one or two oxo or thioxo substituents,

and any of the R<sup>4</sup> groups defined hereinbefore which comprises a CH<sub>2</sub> group which is attached to 2 carbon atoms or a CH<sub>3</sub> group which is attached to a carbon atom may optionally bear on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, amino, C<sub>1-6</sub>alkoxy, *N*-C<sub>1-6</sub>alkylamino, *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino and heterocyclyl;

R<sup>5</sup> is hydrogen, halo, trifluoromethyl, cyano, nitro, amino, hydroxy, C<sub>1-6</sub>alkyl, C<sub>2-6</sub>alkenyl,

C<sub>2-6</sub>alkynyl, C<sub>1-6</sub>alkoxy, *N*-C<sub>1-6</sub>alkylamino or *N,N*-(C<sub>1-6</sub>alkyl)<sub>2</sub>amino;

q is 0, 1, 2, 3 or 4;

or a pharmaceutically acceptable salt or an *in vivo* cleavable ester thereof;

with the proviso that 7-amino-4-(3-acetamidoanilino)pyrido[4,3-*d*]pyrimidine is excluded.

B<sup>2</sup>

10. (Amended) A pharmaceutical composition which comprises a bicyclic compound of the Formula (I), or a pharmaceutically acceptable salt or *in vivo* cleavable ester thereof, according to any one of claims 1-8 in association with a pharmaceutically acceptable diluent or carrier.

B<sup>3</sup>

12. (Amended) A method of treating a disease or medical condition mediated by cytokines which comprises administering to a warm-blooded animal in need thereof an effective amount of a bicyclic compound of the Formula (I), or a pharmaceutically acceptable salt or an *in vivo* cleavable ester thereof, according to any one of claims 1-8.

Please add the following new claim 13:

B<sup>4</sup>

13. (New) A method of treating a disease or medical condition mediated by cytokines which comprises administering to a warm-blooded animal in need thereof an effective amount of the compound 7-amino-4-(3-acetamidoanilino)pyrido[4,3-*d*]pyrimidine.